AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1 (currently amended). A yeast promoter which comprises at least 17 contiguous nucleotides of an An isolated and purified polynucleotide consisting of which is-SEQ ID NO:2, wherein the promoter polynucleotide is operative as a promoter to express a nucleic acid molecule encoding a polypeptide when operably linked to said nucleic acid molecule.

2-8 (cancelled)

9 (currently amended). A yeast expression vector comprising the yeast promoter polynucleotide of claim 1.

10 (currently amended). The yeast expression vector of claim 9 wherein the yeast expression vector is selected from the group consisting of pYLR110P+luc, pYMR251AP+luc, andpYMR107P+luc, pZEO1P+luc, pYLR110P, pYMR251AP, pYMR107P, and pZEO1P.

11-17 (cancelled)

18 (currently amended). A yeast cell transformed with the yeast expression vector of claim 9.

19 (currently amended). A yeast cell transformed with the yeast expression vector of claim 10.

20 (currently amended). A method for producing a polypeptide comprising the steps of:

- (a) constructing a yeast expression vector wherein a nucleic acid encoding the polypeptide is controlled by the <u>yeast promoter polynucleotide</u> of claim 1;
 - (b) transforming a culture of yeast cells with the yeast expression vector;
- (c) maintaining the yeast cells in culture so that the polypeptide is expressed; and
 - (d) recovering the polypeptide.

21 (currently amended). A method for producing a polypeptide comprising the steps of:

- (a) cloning a nucleic acid molecule encoding the polypeptide into an expression vector selected from the group consisting of pYLR110P+luc, pYMR251AP+luc, andpYMR107P+luc, pZEO1P+luc, pYLR110P, pYMR251AP, pYMR107P, and pZEO1P, wherein the nucleic acid molecule is operably linked to a promoter of the expression vector;
 - (b) transforming a culture of yeast cells with the yeast expression vector;
- (c) maintaining the yeast cells in culture so that the polypeptide is expressed; and
 - (d) recovering the polypeptide.

- 22 (currently amended). A method for producing a polypeptide comprising the steps of:
- (a) constructing a yeast expression vector wherein a nucleic acid molecule encoding the polypeptide is controlled by, the polynucleotide of claim 1; a yeast promoter which comprises at least 17 contiguous nucleotides of an isolated and purified polynucleotide which is SEQ ID NO:2;
 - (b) transforming a culture of yeast cells with the yeast expression vector;
- (c) maintaining the yeast cells in culture medium and controlling the expression of the nucleic acid molecule encoding the polypeptide by varying the level of a fermentable carbon source in the culture medium; and
 - (d) recovering the polypeptide.
- 23 (currently amended). The method of claim 22 wherein the fermentable carbon source is glucose.
- 24 (currently amended). A method for producing a polypeptide comprising the steps of:
- (a) constructing a yeast expression vector wherein a nucleic acid molecule encoding the polypeptide is controlled by the yeast promoter polynucleotide of claim 1;
 - (b) transforming a culture of yeast cells with the yeast expression vector;

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- (c) maintaining the yeast cells in culture medium and controlling the expression of the nucleic acid molecule encoding the polypeptide by varying the level of a non-fermentable carbon source in the culture medium; and
 - (d) recovering the polypeptide.

25 (currently amended). The method of claim 24 wherein the non-fermentable carbon source is ethanol.

26 (currently amended). A method for producing a polypeptide comprising the steps of:

- (a) constructing a yeast expression vector wherein a nucleic acid molecule encoding the polypeptide is controlled by the polynucleotide of claim 1;a yeast promoter which comprises at least 17 contiguous nucleotides of an isolated and purified polynucleotide which is SEQ ID NO:2;
 - (b) transforming a culture of yeast cells with the yeast expression vector;
- (c) maintaining the yeast cells in culture medium and controlling the expression of the nucleic acid molecule encoding the polypeptide by varying the level of a fermentable carbon source and a non-fermentable carbon source in the culture medium; and
 - (d) recovering the polypeptide.

27 (currently amended). The method of claim 26 wherein the fermentable carbon source is glucose.

28 (currently amended). The method of claim 26 wherein the non-fermentable carbon source is ethanol.

29 (currently amended). A method of identifying a promoter fragment, wherein the fragment has promoter activity comprising the steps of:

- (a) generating a fragment comprising at least 17 contiguous nucleotides of an isolated and purified polynucleotide which is consisting of SEQ ID NO:2;
- (b) cloning the fragment into a yeast expression vector, wherein the fragment is operably linked to a reporter gene;
 - (c) transforming yeast cells with the yeast expression vector;
- (d) growing the yeast cells in yeast cell culture under conditions favorable for expression of the reporter gene; and
- (e) assaying the yeast culture for a reporter protein expressed by the reporter gene;

wherein expression of the reporter gene indicates the fragment has promoter activity.